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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C08L 23/06, D06M 15/21 C09J 7/02		A1	(11) International Publication Number: WO 92/04409 (43) International Publication Date: 19 March 1992 (19.03.92)
(21) International Application Number: PCT/US91/06342			(74) Agent: ROONEY, Gerard, P.; Allied-Signal Inc., Law Department (C.A. McNally), P.O. Box 2245R, Morristown, NJ 07962-2245 (US).
(22) International Filing Date: 5 September 1991 (05.09.91)			
(30) Priority data: 578,486 7 September 1990 (07.09.90) US 698,230 10 May 1991 (10.05.91) US			(81) Designated States: AT (European patent), BE (European patent), CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, KR, LU (European patent), NL (European patent), SE (European patent).
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(54) Title: COEMULSIFICATION OF OXIDIZED POLYETHYLENE HOMOPOLYMERS AND AMINO FUNCTIONAL SILICONE FLUIDS

(57) Abstract

Articles comprising coemulsions of oxidized polyethylene with amino functional silicone fluids which includes at least one constituent selected from the group consisting of: ethoxylated aliphatic amines, ethoxylated nonylphenols, ethoxylated primary alcohols, ethoxylated secondary alcohols or other nonionic emulsifiers are taught, as well as processes for making the same. The coemulsions may also comprise further constituents, including but not limited to fatty acids, ammonium hydroxide, salts and water. Processes for the production of the coemulsion and articles comprising the same are also disclosed.

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COEMULSIFICATION OF OXIDIZED POLYETHYLENE HOMOPOLYMERS

AND

AMINO FUNCTIONAL SILICONE FLUIDS

BACKGROUND

5 1. Field of the Invention

The invention relates to coemulsions of amino functional silicone fluids and oxidized polyethylene and processes for making the same.

10 2. Description of the Prior Art

Emulsions of various organic compounds are well known in many arts for their widespread utility. Exemplary are the use of emulsions in cosmetic compositions, including creams, lotions, cosmetic 15 compositions, and the like. Other examples include uses as lubricating agents for sheet-like materials, including papers and fabrics, softening agents for textiles, fiber lubricants, paper release agents, metal working, coatings and as oil-recovery fluids from 20 subterranean formations.

One particular use of emulsions comprising silicones and other organic materials, such as polyethylene are in the area of textiles, for such purposes as lubricants which aid during textile 25 processing, and as conditioners to soften textiles or otherwise improve the tactile properties, such as the "hand" of the fabric (the term "fabric" is interchangeably called "textile") are well known. Further benefits of the use of such emulsions as 30 lubricants include reduction of needle cutting and needle abrasion, improved abrasion resistance, increases in tear strength, and cr ase recovery.

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Examples of such known compositions include U.S. Patent No. 4,394,518 to Huber et al. for "Organic Fibers Having Improved Slip Properties" which is directed to a polymeric organosilicon compound 5 containing aryl radicals and methods for using the compound for improving the slip properties of fibers. A further example is taught in U.S. Patent No. 4,767,646 to Cordova et al. for "Wet Abrasion Resistant Yarn and Cordage" which teaches an oxidized 10 polyethylene emulsified with a non-nitrogen nonionic emulsifiers and neutralized with an alkali hydroxide, and which further includes a amide melamine wax and a further siloxane containing compound. U.S. Patent 3,844,826 to Buchner et al. for "Dressing Sewing Thread 15 to Reduced Friction" teaches a sewing thread having improved slip characteristics which includes a non-friction coating of a polycarbonate-polydimethylsiloxane block copolymer, which preferably further includes a polydiorganosiloxane in its 20 composition.

Technical Data sheet titled "The Use of A-C® Polyethylenes in Textiles" describes a family of emulsifiable polyethylenes and methods for making emulsion which include these products, as well as 25 beneficial features which are imparted unto textiles through the use of these materials and compositions. However, this data sheet does not teach or suggest the invention to be presented herein.

While it is known to the art that an emulsion 30 formed as a combination or comixture of an oxidized polyethylene- containing emulsion and a silicone-containing emulsion provide good resultant properties, including good lubricating properties for both threads, fibers and woven fabrics, the relative cost of silicon-containing emulsions curtail their broader use. None 35

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of the art teaches a composition and a process for using a composition wherein an oxidized polyethylene and a silicone containing material are coemulsified and the emulsion so formed is then used to provide improved 5 slip characteristics to a broad range of filament, fibrous, fabric and sheet type materials.

SUMMARY

There are provided articles comprising a 10 coemulsion, wherein the coemulsion is a stable aqueous coemulsion of oxidized polyethylene or polyethylene copolymers with amino functional silicone fluids which includes at least one constituent selected from the group consisting of; ethoxylated aliphatic amines, 15 ethoxylated octylephenols, ethoxylated nonylphenols, ethoxylated primary alcohols, ethoxylated secondary alcohols or other nonionic emulsifiers. The coemulsions may also comprise further constituents, including but not limited to fatty acids, ammonium 20 hydroxide, salts and water.

There is also provided a method of producing a coated article comprising a coemulsion of oxidized polyethylene with amino functional silicone fluids, as described above wherein such method comprises process 25 steps of: heating said constituents in the presence of water with agitation in a sealed and pressurized reaction vessel through the melt point of the oxidized polyethylene constituent, further raising the temperature of the constituents beyond the melt point 30 of the oxidized polyethylene constituent to a maximum temperature, maintaining the constituents at this maximum temperature for a first residence time interval, cooling the constituents, and ultimately contacting an article with the coemulsion produced.

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DESCRIPTION OF THE PREFERRED EMBODIMENTS

Throughout the body of the specification and the claims, all percentages are to be understood as weight percentages relative to a particular overall composition, unless specifically indicated as otherwise.

Suitable polyethylenes may be characterized as oxidized low density and high density homopolymers of ethylene, copolymers containing acrylates and ethylene, terpolymers containing acrylates, esters and ethylene. These polyethylenes have preferably been oxidized to an acid number as determined by a standardized titration of KOH of about 5 and 55, more preferably between about 10 and 50, and most preferably between about 10 and 45.

These polyethylenes typically have a density as determined by ASTM D-1505 in the range of about 0.85 to 1.05, more preferably in the range of about 0.87 to about 1.05, and most preferably, in the range of about 0.90 to about 1.00. Preferably, these oxidized polyethylenes exhibit a Brookfield viscosity at a temperature of 140 deg.C of between about 185 and 6000 centipoises (hereinafter interchangeably referred to as "cps"), more preferably in the range of about 190 and 6000 cps, and most preferably in the range of between about 190 and 5500 cps. Such oxidized polyethylenes are currently commercially available from the Allied-Signal Corp. under the designations of A-C® 629 and 392. The former is described as having a density of 0.93 an acid number of 16 and a Brookfield viscosity of 200 cps, while the latter is described as having a density of 0.99, an acid number of 30 and a Brookfield viscosity of 4500 when measured at a temperature of 4500 deg C.

These oxidized polyethylenes as well as others which are useful in the practice of the instant

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invention may be obtained by oxidizing polyethylenes with air or oxygen by conventional procedures.

Suitable methods are described in U.S. Patent No.

3,060,163 to Erchak, Jr., U.S. Patent No. 3,322,711 to

5 Bush et al., which methods are all hereby incorporated by reference.

Appropriate amino functional silicone fluids which may be used in accordance with the invention are those which may be broadly described as amino organic

10 modified polysiloxanes, alternatively known to the art as "polydiorganosiloxanes". Preferable

polydiorganosiloxanes include those which include the characteristic of exhibiting an amine neutral

equivalent in the range of approximately between about 15 1000 and 3000, more preferably in the range of between about 1200 and 3000, and most preferably in the range of between about 1250 and 2800. Examples of

commercially available polydiorganosiloxanes which find use in the instant invention include those which are 20 marketed under various trade designations. One suitable group of materials include Dow Corning® CSF and Dow Corning® SSF which are sold as textile softening and lubricating compositions. These

materials are described as medium viscosity 25 polydiorganosiloxane which comprise aminoalkyl groups affixed to a predominantly polydimethylsiloxane structure. The amine neutral equivalent is described to be approximately 2000, the specific gravity at 25

deg.C (77 deg.F) of 0.96, and to have a viscosity at 25

30 deg.C (77 deg.F) of 1300 centistokes (hereinafter interchangeably referred to as "cst"). A further commercially available material includes UCARSIL® Magnasoft™ materials available from Union Carbide Corporation and marketed as textile softening 35 compositions. These materials are described to be low

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viscosity amino functional silicones which have a viscosity of about 250 cst at a temperature of 25 deg.C (77 deg.F), a specific gravity of 0.97 at 25 deg.C (77 deg.F). A further commercially available

5 polydiorganosiloxane material includes those which are marketed by PPG-Mazer under the designations MASIL® 123 and MASIL® 124.

As had been noted, the microemulsion includes at least one constituent selected from the group

10 consisting of; ethoxylated aliphatic amines, ethoxylated octylphenols, ethoxylated nonylphenols, ethoxylated primary alcohols, ethoxylated secondary alcohols. The at least one constituent selected from this group which is added to the polyethylene and the

15 amino functional silicone fluid in accordance to the teaching of this present invention may be interchangeably referred to as the "additive system", and it is to be understood that the "additive system" includes only one or more of constituents selected

20 from: ethoxylated aliphatic amines, ethoxylated octylphenols, ethoxylated nonylphenols, ethoxylated primary alcohols, ethoxylated secondary alcohols. It is further to be understood that while further composition, constituents, reagents and the like might 25 find use in conjunction with the present invention, and which might be known to the art as useful additives, the term "additive system" as used throughout this specification and the claims are to be understood to be limited to the group of four constituents described.

30 The ethoxylated aliphatic amines which are suitable to the practice of the invention are those which may be described as saturated and unsaturated fatty amines reacted with ethylene oxide. These materials which are useful in the practice of the 35 present invention may be generally termed as the

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condensation products of ethylene oxide with a hydrophobic material such as a long chain aliphatic alcohol, ester, acid, ether or alkyl phenol. These materials which find use in conjunction with the invention are characterized by containing as the hydrophilic portion of the molecule a plurality of oxyethylene moieties. Suitable materials of this type may also be referred to as ethoxylated tallow amines, a designation commonly used in the art.

Examples of preferred ethoxylated aliphatic amines which are presently commercially available include "Ethomeen T-12" and "Ethomeen 18/12" which is available from Akzo Chemie America and which is described as an ethoxylated tallowamine, more particularly described as bis-(2-hydroxyethyl) tallowamine. A further example includes "Varonic T-202" which may be described as a constituent having similar characteristics to those described in conjunction with Ethomeen T-12 above, and are believed to be functionally identical. Varonic T-202 is at present commercially available from the Sherex Co., Chicago Ill. Further preferred ethoxylated amines include ethoxylated octylphenols and nonylphenols which may be described as being the reaction products of an octylphenol or a nonylphenol with ethylene oxide. Examples of preferred ethoxylated octylphenols and nonylphenols which find use with the instant invention include those which are sold under the designation "Igepal CO-430" which is sold as a surfactant and available from GAF Corporation and which is described as an ethylene oxide, more particularly as nonylphenoxy poly(ethyleneoxy)ethanol having a molecular weight of 484, and a boiling point in excess of 93.30 deg.C.

Preferred thoxylated alcohols include ethoxylated primary alcohols and ethoxylated secondary alcohols

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suitable in the practice of the present invention include those which may be described as the reaction product of a primary alcohol or a secondary alcohol and ethylene oxide. Examples of such commercially

5 available ethoxylated alcohols include; "Tergitol 15-S-3" available from Union Carbide Corp. of Danbury, Connecticut which is described as an ethoxylated secondary alcohol, "Ethal TDA-3" which is described as an ethoxylated tridecyl alcohol formed as the reaction
10 product between stoichiometric quantities of 3 moles of ethylene oxide with one mole of tridecyl alcohol, "Neodol" which is commercially available from Shell Oil Corp.

According to the invention, the polyethylene and
15 amino functional silicone fluid further includes the additive system which consists of at least one constituent selected from among the group consisting of: ethoxylated aliphatic amines, ethoxylated octylphenols, ethoxylated nonylphenols, ethoxylated
20 primary alcohols, ethoxylated secondary alcohols. Where the additive system consists of only one constituent, then it is to be recognized that the selected constituent forms 100 percent of the additive system's composition. It is preferred however that the
25 additive system comprise at least two components where one component is an ethoxylated aliphatic amine, and the other component or components are selected from the remaining members of the group, namely the ethoxylated nonylphenols, ethoxylated primary alcohols, and
30 ethoxylated secondary alcohols. More preferably, the composition of the additive system consists of the following constituents in the following proportions: ethoxylated aliphatic amines 40 - 100 % mol., ethoxylated nonylphenol, 0 - 60 % mol., ethoxylated
35 primary alcohols, 0 - 60 % mol., ethoxylated secondary

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alcohols, 0 - 60 % mol.. Most preferably, the relative molar percentages of the constituents making up the additive system are present in the following ratios: ethoxylated aliphatic amines 60 - 100 % mol.,

5 ethoxylated nonylphenol, 0 - 40 % mol., ethoxylated primary alcohols, 0 - 40 % mol., ethoxylated secondary alcohols, 0 - 40 % mol.

Additional constituents may be incorporated in to the inventive compositions, and such constituents may 10 be in any amount which does not detract from the benefits of the invention.

Water may be present in any quantity which does not have any detrimental effect upon the formation of the emulsion according to the teachings of the 15 invention. Preferably, quantities of between about 30 to 90 percent by weight of the total composition, more preferably between about 45 to 90 percent by weight, most preferably between about 60 to 80 percent by weight of water are included in the total composition.

20 A further constituent which may be used in formulating microemulsion compositions include acetic acid in either glacial or dilute forms, which may be added in order to modify the relative acidity of the composition by adjusting the respective pH value 25 thereof. Preferably, the pH should be maintained to adjust the relative affinity to the ethoxylated amine so that material exhibits a greater affinity to the olefins in the overall composition rather than an affinity to any water contained in the overall 30 composition.

Hydrochloric acid, in either concentrated, i.e. approximately 37% concentrations, or dilute concentrations, i.e., 12% or less, may also be included in formulating the inventive compositions. When added, 35 hydrochloric acid functions to provide a means to

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adjust the relative pH of the overall compositions as is described above.

Ammonium hydroxide is a further useful reagent which may be included in the composition and 5 preferably, concentrated ammonium hydroxide of an approximate 30% concentration may be used. It has been observed that the presence of ammonia, in even minor amounts in microemulsion compositions according to the present invention imparts a beneficial effect upon the 10 composition which is evidenced by reductions in the opacity of the microemulsions formed containing ammonia relative to the opacity of the microemulsions formed without containing ammonia. This is evident from the improved appearance and measured light transmission 15 measurements which are factors known to the art to be indicative of the relative particle size in an emulsion.

Sodium salts, including but not limited to sodium chloride, sodium hydroxide, and sodium metabisulfite 20 may be incorporated as a constituent, as the salt provides the desirable effect of reducing the viscosity, and where the salt is sodium metabisulfite, to improve the color and optical characteristics of the composition. The sodium salts may be in any form, such 25 as a finely divided powder, or in a pelletized form, or alternately, the sodium salt may be dissolved in a reagent or other constituent, for example in the water comprising a composition which is subsequently combined with the other constituents. The salts may also be in 30 various percentages, and may further include small amounts, i.e., 20 % or less of inert materials which exhibit no detrimental effect upon the practice of the invention.

It is recognized and understood that although only 35 particular constituents have been recited above, other

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materials which are known to the art which find use in the compositions of emulsions may be included, and such additives are considered as part of the invention disclosed herein.

5 The process for forming the microemulsions of the present invention are formed include the following process steps: heating the constituents used to form the microemulsion under conditions of constant agitation in a sealed and pressurized reaction vessel

10 through the melt point of the oxidized polyethylene constituent, further raising the temperature of the constituents beyond the melt point of the oxidized polyethylene constituent to a maximum temperature, maintaining the constituents at this maximum

15 temperature for a first residence time interval, then cooling the constituents. In an alternative embodiment of the process, the heating of the constituents comprises two heating steps; applying heat at a first heating rate to a first intermediate temperature and

20 then heating the constituents from the first intermediate temperature to the maximum temperature at a second heating rate. Preferably, the first heating rate is greater than the second heating rate, i.e., a faster transmission of heat per unit of time occurs

25 during the first heating rate than during the second.

Any suitable vessel capable of withstanding the operating parameters of the process may be used in conjunction with the present invention. By way of example, suitable reactors include those constructed of

30 metals, glass, glass-lined metal reactors. Preferably, the reaction vessel is a batch type reaction vessel which may be sealed, and which is of sufficient strength so to withstand the pressure generated by the constituents during the process of forming the

35 microemulsion.

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It is contemplated that the reaction may be carried out in any manner, either in a batch type mode, or in a continuous mode or other mode of operation; any reaction mode which is successful in providing the 5 conditions for the productions of the emulsions taught herein may be utilized. Preferably the reaction is carried out in a batch type mode.

Agitation means are meant to include any means, or alternately any method by which the constituents in the 10 reactor may be well mixed, and by way of example, such means include rotary paddles, propellers, stirring rods, vanes and the like, as well as other methods which may comprise shaking, spinning or otherwise effecting movement of constituents within the reactor.

15 In a preferred embodiment of the process the selected constituents comprising an oxidized polyethylene, amino functional silicone fluid, and the additive system as defined above, as well as other desired constituents are loaded into a reaction vessel 20 which is sealed, and which includes agitation means. The selected constituents are heated, with agitation, at a first heating rate through the melting point of the oxidized polyethylene component for a first time interval, and then during a second time interval 25 further heated at a second heating rate to raise the temperature to a maximum temperature or maximum temperature range approximately 7-10 deg.C beyond the melt point of the oxidized polyethylene component, and then maintaining the reactants at this maximum elevated 30 temperature or maximum temperature range for a period of between about 10-15 minutes. After the period had elapsed, the materials were then cooled to room temperature. Agitation is maintained throughout the procedure.

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The order of adding or combining the constituents are not critical, and they may be added in any order as is suitable for use with the reactor and the reaction method employed.

5 The relative acidity of the composition, i.e., the "pH" should be evaluated in order assure that the microemulsion is of suitable optical clarity. Preferably, the pH may be adjusted by modifying the amounts of the constituents used, especially by
10 modifying the amount of acetic acid and/or hydrochloric acid used. Preferably, the pH of the microemulsion should be maintained in a range of between about 4 to 6, more preferably in the range of between about 4.5 and 5.5, and most preferably in the range of between
15 about 4.75 and 5.25.

The coemulsification products of this procedure form stable coemulsions of the oxidized polyethylene and amino functional silicone fluid, which include particles in the size range of between about 0.0001
20 and 100 microns. Preferably, the products of this procedure have particle sizes in the range of between about 0.001 and 5 microns, and most preferably comprise particles having particle sizes in the range of about 0.001 and 1 micron. The small particle sizes of the
25 present invention are generally classed as characteristic of a "microemulsion". As is well known in the art, the smaller particle sizes of the emulsion enhance the distribution of the silicon in the emulsions, and improve the distribution of the silicon
30 within or upon the surface of any material, such as textile, paper or other sheet type material and thereby impart such desirable properties as lubrication, textural modification or other property with a consequent reduction in the amount of silicon or other
35 constituent required to form a suitable coemulsion.

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Determination of the optical characteristics of each of the compositions, as well as any control compositions were evaluated to determine their optical density from which the optical transmittance, and 5 conversely, relative opacity, of a composition may be derived. This procedure is well known to the art to be a useful index of the average particle size contained in compositions. The evaluation of optical density was performed with the use of a Klett-Summerson 10 Photoelectric Colorimeter, Model No. 800-3, which was equipped using a number 66 filter which is rated to have a spectral range of between 640 and 700 millimicrons. The values provided by the Klett-Summerson Photoelectric Colorimeter are termed "Klett 15 Values" from which a corresponding optical density and a percent light transmittance can be determined.

A further optical evaluation of the inventive coemulsions and a blend of two emulsions, i.e., a polyethylene emulsion and an amino silicone containing 20 emulsion was performed on a Leitz Ortholux optical microscope. The magnification was varied as described below, and a xenon lamp lighting source was used in either of two lighting modes, phase contrast or what is commonly referred to in the art as a "transmitted dark-field" in order to improve the contrast of the 25 particular samples. The results of this evaluation is shown on Figures 1-8. Figures 1-3 illustrate a cationic emulsion formed as a result of a blend of two emulsions produced by producing a mixture under 30 pressure at a temperature in the range of 110 - 115 deg.C of a low density oxidized polyethylene and 2 moles of an ethoxylated tallowamine surfactant in a reactor with stirring, using Masil® EM 115, a 35 polydimethylsiloxane material commercially available from PPG-Mazer chemicals. The two blended emulsions

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were observed at magnifications of 260x, 650x and 1040x. Figures 4-6 illustrate a coemulsion formed in accordance with the teaching of the instant invention, whose composition is described below as Example K1.

5 Similarly, Figures 7-8 illustrate an inventive coemulsion whose composition is designated below as Example C14. As may be seen by inspection, Fig. 1 which was produced using dark-field illumination illustrates particles which are generally non-uniform 10 in geometrical configuration, particularly the particles denoted as 1,2 and 3. These particles have an amorphous and irregular configuration. Turning now to Fig.2, illuminated using phase contrast lighting the same sample is viewed under a higher magnification, and 15 two distinct particles are particularly apparent; the particle denoted as 4 is seen to have an irregular shape a particles illustrated in conjunction with Fig.1, and the particle denoted 5 is seen to have a spherical or circular shape. The following Fig.3 20 illuminated by phase contrast lighting, shows the same sample as the two prior magnifications at a higher magnification, and the particle denoted as 6 reveals a generally spherical particle, which is approximately 10 microns in diameter. The hazy interior suggests that 25 the particle 6 has no internal structure, and it is hypothesized that the particle 6 is a suspended oil particle.

Turning to Figures 4-6, there is illustrated a sample of the coemulsion formed in accordance with the 30 teachings of the present invention having the composition of the coemulsion labeled "K1" of Table K-1 noted below. Using a dark-field lighting mode, wherein the light from the light source passes at an angle nearly oblique to the sample, Figure 4 illustrates a 35 more uniform distribution of the particles resulting

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from the coemulsion technique taught in the instant specification. As may be seen thereon, and in contrast to the sample illustrated on Figures 1-3, all of the particles distinctly visible on Figures 4-6 are of a

5 generally spherical configuration, and the particles contain an internal structure. Such structure is most clearly seen on Figure 5 which is also illuminated under a dark-field lighting mode, where a particle denoted as 7, an less readily visible from Figure 4 in

10 particles labeled 8 and 9. Upon inspection, an internal structure is visible within these particles, especially when viewed in contrast to Figures 1-3.

While not fully understood, and not wishing to be limited to any particular theory, it is hypothesized

15 that the structures observed in Figures 4-8 reveal an internal structure which comprises small pieces of polyethylene intermixed in silicone oil which suggests the formation of coemulsified particles of the polyethylene and the silicone rather than separate

20 particles of polyethylene and separate droplets and/or particles of silicone, which is suggested as descriptive of the particles illustrated in Figures 1-3.

Figure 6, illuminated under phase contrast lighting

25 further shows visible particles having the internal structure noted above. Further, Figures 7 and 8 both illuminated under dark-field lighting modes illustrate two further coemulsions according to the compositions of samples C14 as listed on Table C1 noted below. It

30 should be noted that Figures 7 and 8 further illustrate successful coemulsions which reveal particles having internal structures apparently identical in nature to the particles illustrated on Fig. 4-6.

The samples utilized for the production of Figs.

35 1-8 were further analyzed by photon correlation

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spectroscopy by methods which are outlined in the following references: *Laser Light Scattering*, by B. Chu published by Academic Press, New York, 1974; *Dynamic Light Scattering*, by B. Berne and R. Pecora, published 5 by Wiley, New York, 1976 and *Dynamic Light Scattering: Applications of Photon Correlation Spectroscopy*, R. Pecora, editor published by Plenum Press, New York, 1985. Determination of the size distribution of particles may be determined from photon correlation 10 analysis by methods which are outlined in the following references: Provencher, S.; Hendrix, J.; DeMaeyer, J. *J.Chem.Phys.*, 1975, Vol. 69, p. 4273 and Provencher, S. *Makromol. Chem.*, 1979, Vol. 180, p.201. The samples of Figs 1-8 were analyzed in accordance with the methods 15 described in the publications cited, the contents of which are herein incorporated by reference, and weight-average particle diameter "d_w", number-average particle diameter "d_n" and z-average particle diameter, "d_z" were determined. These values are defined in 20 accordance with the following formulas:

$$d_w = \frac{\sum_i^N n_i M_i d_i}{\sum_i^N n_i M_i}$$

$$d_n = \frac{\sum_i^N n_i d_i}{\sum_i^N n_i}$$

$$d_z = \frac{\sum_i^N n_i M_i d_i^2}{\sum_i^N n_i M_i}$$

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$$d_z = \frac{\sum_i^N n_i M_i^2 d_i}{\sum_i^N n_i M_i^2}$$

10 where, M_i is representative of the mass of a particle,
 15 d_i is representative of the diameter of a particle, and
 n_i is the number of particles having a unit mass. The
 20 diameters in units of microns (μm) are listed in Table
 25 PD below.

TABLE PD

20	Sample Evaluated:	<u>Resultant Diameters, (μm)</u>		
		d_w	d_n	d_z
	Figs. 1-3	0.028	0.007	0.145
25	Figs. 4-6	0.023	0.006	0.076
	Figs. 7-8	0.012	0.005	0.069

30 From these reported results, it can be seen that while
 35 there was no distinctive differences in the number-average diameter, d_n of the particles, and moderate differences in the weight-average diameter, d_w , of the
 40 particles, between the samples formed from the coemulsions in according with the teachings of the present invention (the samples of Figs. 4-6 and Figs. 7-8), there is a striking difference in the evaluated z-average particle diameter, d_z , which, for the inventive coemulsions has been observed to be less than for blended emulsions. This figure suggests that the average diameter of the coemulsion particles are

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generally smaller than that for blended emulsions, and while not wishing to be bound to this theory, might further be suggestive that they would provide an improvement in lubricative characteristics.

5 The coemulsions described above may be contacted to a variety of articles so to provide one or more of the following characteristic features: improved lubricity, improved anti-adhesive properties, improved tactile properties particularly for fibers, threads and
10 textile applications, improved mold release agents for dies and molds for forming articles from thermoplastic resins.

15 Articles may be imparted with improved lubricative characteristics by contacting the articles with coemulsions described above by any effective method which method includes conventional methods, which methods include but are not limited to: immersion of the article, immersion of at least part of the article, spraying, coating, brushing-on, or otherwise contacting
20 the coemulsions with the article.

25 Articles which are wholly or at least partially comprised of fibers including but not limited to textiles, fibers, ropes, tows, webs, threads, so-called "fiberfill" materials, as well as others are imparted with improved lubricity, improvements in the tactile quality of the article as is commonly referred to in the textile art as "hand", as well as improved softness of the article.

30 Paper and other flexible sheet type articles including but not limited to paper sheets, sheets and films formed from of thermoplastic compositions such as polyesters, may be provided with an anti-adhesive coating and with improved slip characteristics. Paper and flexible sheet type articles coated with the
35 coemulsion of the present invention provide a useful

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carrier or release coating for a wide range of adhesives, and other materials which exhibit adhesive properties.

One class of adhesives with which the coemulsions of the present inventions find use are with pressure sensitive adhesives and other adhesives which are desirably applied to the surface of one material, upon to which a release paper, release strip or release sheet is applied so to protect the pressure sensitive adhesive until ready for use. In use, the release paper, release strip or release sheet is removed, leaving a major portion of the pressure sensitive adhesive in contact with the material upon which it was applied. Examples of such articles include tapes, sheets, decals, labels, envelopes, as well as other paper and sheet type articles as well as other articles not enumerated here upon which the pressure sensitive adhesive may be applied. The coemulsions of the present invention find particular use as a coating for the release sheet, release film, release strip or release paper described and feature poor adhesion to the pressure sensitive adhesive, which is desirable property.

A further class of materials which exhibit adhesive properties include asphalt containing compositions and articles comprising asphalt. Examples of such articles include roofing shingles, and other roofing materials, and packaging containers for asphalt containing compositions. The coemulsion compositions taught in the instant invention provide structures featuring good release properties for use in conjunction with asphalt containing compositions and articles comprising asphalt.

Roofing shingles such as those presently in common use in North America for buildings, residential

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buildings and other structures may be generally described as a substantially sheet type article having a length dimension greater than a width dimension, and two opposite faces, an "upper face" upon which is 5 conventionally placed a series or alternately a strip of a heat curing adhesive, and on the "lower face" (the reverse side of the roofing shingle) there is desirably placed a release material which is non-adhesive, or exhibits only limited adhesive characteristics when 10 contacted with the heat curing adhesive. During the production, shipment and storage of such roofing shingles prior to their installation of a structure, packaged roofing shingles are layered in register to form a stack wherein the heat curing adhesive of the 15 upper face of a shingle will be retained in contact with the release material on the lower face of an adjacent roofing shingle. It is highly undesirable that, prior to installation of the shingle, that adhesion of adjacent shingles be realized. Presently, 20 it is known to the art to utilize a silicon coated thermoplastic tape, typically comprising poly(ethylene terephthalate) which is applied to the lower face of a roofing shingle. However effective, this solution is a cost and labor intensive solution.

25 The coemulsions of the present invention when contacted with the lower face of the shingle, preferably in the region of the lower face of the shingle which will contact the heat sensitive adhesive of an adjacent roofing shingle prior to the 30 installation of the roofing shingles provides a very satisfactory release coating which may be directly applied to the lower surface of a roofing shingle and form a layer of the coemulsion thereon. Alternatively, the coemulsions of the present invention may be used to 35 produce a release tape or material of a fiber, paper or

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thermoplastic material such as a polyester tape which is coated with the coemulsion and afterwards applied to the lower face of the roofing shingle in accordance with conventional methods known to the art.

5 Preferably, such a release tape or material for use with a roofing shingle is a paper tape or a sheet, or a polyester tape or sheet; most preferably such a release tape or material is a polyester tape or sheet such as poly(ethylene terephthalate).

10 Packaging containers for asphalt, commonly paper containers which are used to contain a mass of asphalt may be coated on the surfaces of the container which are to contact the asphalt with the coemulsions of the present invention. The coemulsions of the present 15 invention provide improved release properties as compared to an uncoated packaging container.

Generally, it may be broadly stated that the coemulsions taught in the present invention may be substituted in applications wherein it is presently 20 known to the art to use a silicone coating and to replace the silicone coating being used.

The foregoing invention will be more apparent by reference to specific embodiments which are representative of the invention. It is nonetheless to 25 be understood that the particular embodiments described herein are provided for the purpose of illustration, and not be means of limitation, and that it is to be further understood that the present invention may be practiced in a manner which is not exemplified herein 30 without departing from its scope.

EXAMPLES

In the descriptions of the following compositions, as well as the tables, all measurements relating to the 35 individual constituents are listed in units of grams

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unless indicated otherwise. The mulision, was evaluated by immersing a portion of a clean glass laboratory slide into the reaction product and withdrawing it, at which point the appearance of the 5 slide was observed.

Examples C1-C14

In the formulations for the compositions according to Examples C1-C14 a stainless steel reactor having a 10 capacity of 2 liters and equipped with an electrical resistance-type heating system in the form of coils which was placeable near the exterior of the reactor, and a stirrer attached to variable speed electric motor, as well as a thermometer. This reactor is 15 interchangeably referred to as the "reactor", "reaction vessel" or "vessel". This stainless steel reactor was emptied and thoroughly cleaned before the formulation and production of each of the Examples unless otherwise specifically indicated.

20 The resultant reaction products of each of the compositions, i.e., the emulsions, were evaluated for their Klett values in accordance with the directions of the manufacturer of the Klett-Summerson Photoelectric Colorimeter from which the optical density and percent 25 light transmittance was derived. The relative acidity, or pH values was determined by the use of Corning pH meter equipped with a glass electrode.

Example C1

30 All the constituents according to those outlined in Table C1 except for the 0.5g of sodium chloride were charged into the reactor after which the charged reactor was sealed.

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TABLE C1

Material:	C1	C2	C3	C4	C5	C6	C7	C8	C9
A-C 629	128	128	128	128	128	128*	128*	128*	128*
A-C 316									
Dow CSF	32	32	32	32	32	32	32	32	32
Ethomeen 18/12									
Ethomeen T-12	56.8	56.8	56.8	56.8	56.8	56.8	51.2	51.2	51.2
Igepal CO-430	14.2	14.2	14.2	14.2	14.2	15	12.8	12.8	12.8
Varonic T-202									
glacial acetic acid	8	7.4	7.4	6.4	6.7	7	7.2	7.2	7.2
hydrochloric acid, 10%				5	5	1	2	3.2	1
hydrochloric acid, 37%									
ammon. hydrox., conc. 30%	0.5	0.5	0.5	0.5	0.5	0.2	0.3	0.3	0.3
sodium chloride									
sodium hydroxide									
sodium metabisulfite									
potassium hydroxide									
water	514.4	514.4	514.4	514.4	514.4	514.4	514.4	514.4	514.4
composition pH	5.3	5.3	4.9	4.65	4.85	5.2	5.15	5.05	5.05
Klett value:	425	325	350	—	—	450	400	400	430
optical density	0.85	0.65	0.699	—	—	0.9	0.8	0.8	0.86
light trans., %	14	20.4	20	—	—	12.59	15.86	15.86	13.72
slide appearance	clear	clear	milky	milky	clear	clear	clear	clear	clear

Notes:

- *** = Indicates an acid value of 16.2
- *** = Indicates an acid value of 15.9
- ** + = Indicates an acid value of 16.9
- ** + + = Indicates an acid value of 15.7

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C10	C11	C12	C13	C14	C15	C16	C17	C18	C19	C20	C21
128 *	128 *	128 *	128 **	128 **	128 **	128	128	128	128	128	128 *+
32	32	32	32	32	32	32	32	32	32	32	32
51.2	51.2	51.2	51.2	51.2	51.2	51.2	51.2	51.2	51.2	51.2	51.2
12.8	12.8	12.8	12.8	12.8	12.8	12.8	12.8	12.8	12.8	12.8	12.8
7.5	7.5	7.5	7.5	7.5	7.5	7.5	7	6.5	7.5	7.5	7.5
0.25	0.25	0.25	0.2	0.2	0.2	0.2	0.2	0.2	0.3	0.3	0.2
0.5	0.4	0.5	0.5	0.5	0.5	0.5	0.25	0.25	0.5	0.5	0.5
514.4	514.4	514.4	514.4	514.4	514.4	514.4	515	515	514.4	514.4	514.4
4.95	4.8	5.05	5.05	5.05	5.05	5	5	5.2	5.15	5.1	4.8
242	425	198	340	82	85	450	360	350	118	185	67
0.495	0.85	0.396	0.68	0.164	0.17	0.9	0.72	0.69	0.238	0.38	0.134
32.84	14	40.16	20.9	68.57	67.66	12.6	19	20	58	42.7	73.5
clear	clear	clear	clear	clear	clear	good	clear	clear	clear	clear	clear

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C22	C23	C24	C25	C26
128 *+	144 *+	128 *+	160 *+	160 *+
32	32	32	40	40
51.2	51.2	51.2		
12.8	12.8	12.8	12.8	
7.5	7.5	7.5	51.2	
0.2	0.2	0.2	9.5	
0.5	0.5	0.5	2.5	2.5
514.4	514.4	514.4	514.4	
—	—	—	—	5
—	—	83	—	600
—	—	0.166	—	1.2
—	—	68.3	—	6.33
cloudy	cloudy	clear	—	good

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No pressure or gas was added to the reactor during the production process. The electric motor was then energized so that the stirrer assured that the constituents were well agitated and the contents of the

- 5 vessel were slowly heated at a rate of approximately 2 to 5 deg.C/min until the contents of the reactor reached a temperature of 110 deg.C. The heating system was then stopped, and the reactor and its contents cooled for approximately 15-20 minutes by withdrawing
- 10 the reactor from the heating coils and immersing the reactor in water at a temperature of approximately 20-22 deg.C. During this cooling phase, the stirrer continued to operate so to assure that the contents of the reactor remained thoroughly agitated. When the
- 15 reactor contents reached a temperature of approximately 30-35 deg.C, the stirrer was halted, and the reactor vessel was opened to allow for the addition of the 0.5g sodium chloride in the form of a finely divided powder. The reactor was again sealed, and the operation of the
- 20 stirrer re-initiated. As before, the heating system was initiated and the reactor and its contents were heated at a rate of approximately 2-5 deg.C/min until the constituents in the reactor reached a temperature of 110 deg.C. With the agitation of the constituents
- 25 continuing, the heating system was disengaged and the reactor allowed to cool in a water bath as described before until it reached a temperature of 30-35 deg.C.

Upon evaluation, the Klett value was found to be 425, which corresponds to an optical density of 0.85, 30 and a light transmission of 14%. The relative acidity testing yielded a pH value of 5.3. The glass slide test revealed an emulsion which had a clear appearance. These results are summarized on Table C1.

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Example C2

The reactor vessel as described above used for the production of Example C1 was charged with all of the constituents and sealed. Agitation was initiated and 5 maintained to assure the adequate mixing of the constituents, and the reactor vessel was heated. The temperature of the constituents was raised according to the manner outlined for Example C1. After reaching the temperature of 110 deg.C, heat was supplied to the 10 reaction vessel so to maintain a constant temperature of 110 deg.C for a period of ten minutes. Subsequently, the heat source was deactivated, and the reactor vessel was allowed to cool to a temperature of 15 30-35 deg.C. The agitator motion was arrested, and the vessel opened.

In accordance with the methods described above and those used for Example C1, the emulsion was evaluated and the results are described in Table C1.

20

Example C3

To the reaction product according to Example C2 was added a quantity of dilute hydrochloric acid at an approximate concentration of 10% to adjust the pH of the emulsion to have a value of 4.9 and to evaluate any 25 change in the optical and physical qualities of the emulsion. These test results are outlined on Table C1.

Example C4

The reactor vessel was charged with Varonic T-202, 30 Igepal CO-430, acetic acid, sodium chloride, and water in the proportions outlined in Table C1. The reactor vessel was sealed and the agitator was energized to assure thorough mixing, and the reaction vessel was heated to 25 deg.C, and at a constant temperature of 25 35 deg.C, the contents stirred for a period of 15 minutes.

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Afterwards, the reactor vessel was opened, and approximately 5 milliliters (5 ml) of dilute (approx. 10% conc.) hydrochloric acid was added to adjust the pH from a value of 5.7 to a value of 4.75. Afterwards,

- 5 the amounts of Dow Corning® CSF and A-C®-629 were added, the reactor vessel sealed and repressurized, agitation reinstated and the vessel was heated at a rate of 1 deg.C per minute from an initial starting temperature of 90 deg.C to 110 deg.C, and upon reaching
- 10 110 deg.C, the temperature was maintained within the range of 110 - 111 deg.C for a period of 15 minutes. Subsequently, heat was terminated and the reactor and its contents were cooled to a temperature of 30-35 deg.C. The reactor vessel was opened and the reaction
- 15 products were tested. The results are indicated on Table C1.

Example C5

To the emptied and cleaned reaction vessel were introduced the quantities of Ethomeen T-12, Igepal CO-430, acetic acid and water as indicated on Table C1. The reactor was sealed and the agitator activated to assure thorough stirring. Heat was supplied to the reactor and its contents to raise the temperature to 25 100 deg.C, after which the heat was removed and the reactor was allowed to cool. The agitator was halted, and the vessel opened in order to add approximately 5.4 grams of concentrated hydrochloric acid so to adjust the acidity of the reaction mixture from a pH of 5.7 to 30 a pH of 4.95. Subsequently, the quantities of A-C® 629 and the Dow Corning® CSF were added to the reaction mixture, the reactor then sealed and repressurized and the agitator restarted. The reactor was then reheated to 110 deg.C from an initial starting temperature of 35 approximately 90 deg.C at a rate of temperature rise of

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1 deg.C per minute of time. Upon reaching 110 deg.C, the reactor was retained at this constant temperature for a time interval of 1 minute, after which the heat source was removed, and the reactor allowed to cool.

5 Upon reaching a temperature of approximately 30-35 deg.C, the agitator was halted, and the reactor was opened. The resulting reactant mixture was observed to produce a slightly milky film when tested upon a glass laboratory slide. To the reaction mixture was added

10 0.25 grams of the sodium metabisulfite, after which the reactor was sealed, repressurized, reagitated and reheated to raise the temperature to 115 deg.C. Upon reaching this temperature the heat was removed, and the reactor was allowed to cool to approximately 30-35

15 deg.C. The agitator was halted, and the reactor was opened and to the reaction mixture was added an additional 0.5 grams of sodium metabisulfite, after which the reactor was sealed, repressurized, reagitated and reheated to raise the temperature to 115 deg.C,

20 upon which the heat was removed and the reactor allowed to cool to a temperature of approximately 30-35 deg.C. The resultant emulsion was tested and found to have a pH value of 4.85, and to be milky in appearance.

25

Example C6

To the reactor was added all of the individual constituents as outlined in Table C1 except for the sodium hydroxide. The reactor was sealed and the electric motor was energized in order to drive the

30 agitator. The contents of the reactor were stirred for one hour (1 hr) at a constant temperature of 25 deg.C. Subsequently, heat was supplied to the reactor contents to raise the temperature to 90-95 deg.C at a first rate of approximately 2 deg.C per minute, and then the heat

35 was reduced to a second rate of approximately 1 deg.C

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per minute to raise the reactor contents from 90-95 deg.C to 110 deg.C. Then, the heat was limited to maintain the reactor contents within a temperature range of 110 - 112 deg.C for a period of ten minutes 5 (10 min), after which the heat was removed, and the contents cooled to a temperature of approximately 30-35 deg.C. The agitator was then halted, and the reactor was opened, at which point was added 0.2 grams of sodium hydroxide in the form of pellets of 90% 10 concentration to the reaction mixture, after which the reactor was sealed, repressurized, reagitated and reheated to raise the temperature to 110 deg.C, upon which the heat was once again removed and the reactor allowed to cool to a temperature of approximately 30-35 15 deg.C. The resultant emulsion was tested with the results indicated on Table C1.

Example C7

To the reactor vessel was introduced the AC-629, 20 Igepal CO-430, and sodium hydroxide in the proportions outlined in Table C1, to which was added 304.4 ml of water. Afterwards was added to the reactor vessel a mixture of the acetic acid, and 1 gram of concentrated hydrochloric acid, (37% conc.) in 210 ml of water. 25 The reactor was then sealed and the agitator activated to assure thorough stirring. Heat was supplied to the reactor and its contents to raise the temperature to 105 - 110 deg.C, and maintained for ten minutes within that temperature range, after which the heat was 30 removed and the reactor was allowed to cool. The heat was then terminated and the reactor and its contents were cooled to a temperature of 30-35 deg.C. The reactor vessel was opened and the reaction products were tested, with the results indicated on Table C1.

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Example C8

To the reaction mixture according to Example C7 was further added an additional 1.0 grams of concentrated hydrochloric acid, (37% conc.). The 5 reactor was then sealed and the agitator activated to assure thorough stirring. Heat was supplied to the reactor and its contents to raise the temperature to 110 deg.C, and then maintained for one minute within that temperature range, after which the heat was 10 removed and the reactor was allowed to cool. The heat was then terminated and the reactor and its contents were cooled to a temperature of 30-35 deg.C. The reactor vessel was opened and the reaction products were tested, with the results indicated on Table C1.

15

Example C9

To the reaction mixture according to Example C8 was further added an additional 1.2 grams of concentrated hydrochloric acid, (37% conc.). The 20 reactor was then sealed and the agitator activated to assure thorough stirring. Heat was supplied to the reactor and its contents to raise the temperature to 110 deg.C, and then maintained for one minute within that temperature range, after which the heat was 25 removed and the reactor was allowed to cool. The heat was then terminated and the reactor and its contents were cooled to a temperature of 30-35 deg.C. The reactor vessel was opened and the reaction products were tested, with the results indicated on Table C1.

30

Example C10

The constituents listed on Table C1 were introduced to the reaction vessel, and the vessel was subsequently sealed and the agitator initiated.

35 Subsequently, the temperature of the vessel was raised

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from 22 deg.C to 75 deg.C at a first rate of 2-5 deg.C per minute, and then from 75 deg.C at a second rate of 1 deg.C per minute, to a temperature within the range of 104 - 108 deg.C. Thereafter, the temperature of the reactor contents was maintained in the range of 104 - 108 deg.C for a period of ten minutes (10 min.), after which time heating of the reactor was terminated and the reactor allowed to cool to 30-35 deg.C. The emulsion was evaluated and result are listed on Table C1.

Example C11

The constituents as listed on Table C1 were charged into the reaction vessel, after which the vessel was subsequently sealed and the agitator activated. Subsequently, the temperature of the vessel was raised from approximately 20 deg.C to a temperature range of 78 - 80 deg.C at a first rate of 2-4 deg.C per minute, and then maintained from 75 deg.C within the temperature range of 78 - 80 deg.C for a period of approximately 45 minutes. Afterwards, the temperature of the reaction mixture was further elevated at a second rate of 1 deg.C per minute, to a temperature within the range of 104 - 108 deg.C. Thereafter, the temperature of the reactor contents was maintained in the range of 104 - 108 deg.C for a period of ten minutes (10 min.), after which time heating of the reactor was terminated and the reactor allowed to cool to 30-35 deg.C. The emulsion was evaluated and result are listed on Table C1, with higher Klett values being observed.

Example C12

To the reactor was added all of the individual constituents, including an A-C® 629 constituent having

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an acid value of 16.2 as outlined in Table C1, after which the reactor was sealed and the electric motor was energized in order to drive the agitator. The contents of the reactor were then rapidly heated to 78 deg.C at 5 a first rate of approximately 2 deg.C per minute, and then the heat was reduced to a second rate of approximately 1 deg.C per minute to raise the reactor contents from 78 deg.C to a temperature in the range of 104 - 108 deg.C. Then, the heat was controlled to 10 maintain the reactor within a temperature range of 104 - 108 deg.C for a period of ten minutes (10 min), after which the heat was removed, and the contents cooled to a temperature of approximately 30-35 deg.C. The agitator was then stopped, and the reactor was 15 opened. The resultant emulsion was tested with the results indicated on Table C1.

Example C13

The procedure used in Example C12 was replicated, 20 with the substitution of one constituent, the A-C® 629 with an acid value of 15.9. The resultant emulsion was tested with the results indicated on Table C1. An increase in the Klett value was observed.

25

Example C14

To the reactor was added all of the individual constituents as outlined in Table C1, which included a quantity of ammonium hydroxide. Subsequently, the reactor was sealed and the electric motor was energized 30 in order to drive the agitator. The contents of the reactor were stirred for fifteen minutes (15 min) at a constant temperature of 27 deg.C. Subsequently, heat was supplied to the reactor contents to raise the temperature to 75 deg.C within a first time interval of 35 twenty minutes (20 min.) which was an first average

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rate of temperature rise of 2.4 deg.C per minute. Afterwards, the rate of heating was reduced, and the temperature of the reactor was allowed to rise for a further time interval of approximately thirty-three 5 minutes (33 min.) which was a second average rate of temperature rise of 1 deg.C per minute. When the reactor contents reached 104 deg.C, the heat was reduced and controlled so to maintain the reactor contents within a temperature range of 104 - 109 deg.C 10 for a period of ten minutes (10 min), after which the heat was removed, and the contents cooled rapidly by immersion into a water bath at a temperature of 20-25 deg.C to a temperature of approximately 30-35 deg.C. The resultant emulsion was tested with the results 15 indicated on Table C1.

Example C15

The procedure used in Example C14 was replicated, using the constituents outlined in Table C1. The 20 resultant emulsion was tested with the results indicated on Table C1. A slight increase in the Klett value was observed.

Example C16

25 In this example, the coemulsification of a high density oxidized polyethylene and an aminosilicone oil was performed. The constituents denoted on Table C1 were introduced into the reactor, which was sealed and the constituents stirred at a temperature of 25 deg.C 30 for a period of one hour. The temperature of the constituents was raised in accordance with the following temperature rise profile: at initialization of heating, 25 deg.C; at 30 minutes, 100 deg.C; at 45 minutes, 124 deg.C; at 55 minutes, 129 deg.C; at 60 35 minutes, 133 deg.C; at 62 minutes, 135 deg.C; at 72

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minutes, 138 deg.C, at which point the heat was removed, and the reactor cooled by immersion in a water bath at room temperature. When cooled, the reactor was opened and a glass laboratory slide was immersed 5 into the reaction mixture. The reaction mixture appeared to be clear, with moderately sized particles therein. The reactor was resealed, and the reaction mixture reheated in accordance with the following profile: at initiation of heating, 29 deg.C; at 20 10 minutes, 115 deg.C; at 40 minutes, 136 deg.C; at 44 minutes, 141 deg.C; at 49 minutes, 141 deg.C; upon which the heat was removed and the reactor cooled in a water bath. The product coemulsion was evaluated with the results are denoted on Table C1.

15

Example C17

A further coemulsification of a high density oxidized polyethylene in combination with an amino-silicone oil was produced. The constituents denoted on 20 Table C1 were placed into the reactor which was sealed, stirred and heated in accordance with the following process steps. The constituents were stirred for 45 minutes at a temperature of 27-28 deg.C and then heated in accordance with the following profile: At the 25 initiation of heating, the temperature was 27 deg.C; at 35 minutes, 112 deg.C; at 69 minutes, 130 deg.C; at 76 minutes, 141 deg.C; at 80 minutes, 145 deg.C; at 86 minutes, 144 deg.C, at which point the heat was removed and the reactor cooled in a water bath. The resultant 30 coemulsion was evaluated and the results are listed on Table C1.

Example C18

A yet further coemulsification of a high density 35 oxidized poly thylene in combination with an amino-

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silicone oil was produced. The constituents denoted on Table C1 were placed into the reactor which was sealed, stirred and heated in accordance with the following process steps. The constituents were first stirred

5 within the sealed reactor for 10 minutes at a temperature of 24-29 deg.C and then heated in accordance with the following profile: At the initiation of heating, the temperature was 29 deg.C; at 27 minutes, 100 deg.C; at 45 minutes, 120 deg.C; at 54
10 minutes, 130 deg.C; at 64 minutes, 141 deg.C; at 74 minutes, 144 deg.C, at which point the heat was removed and the reactor cooled in a room temperature water bath. The resultant coemulsion was evaluated and found to be good, and the results are listed on Table C1.

15

Example C19

A coemulsification of a low density polyethylene and having higher acid values than used in prior Examples was formed. The constituents noted on Table C1 were introduced into the reactor which sealed and the reactants stirred with the constituents remaining at 26 deg.C over a period of 15 minutes. The reactor was then heated in accordance with the following temperature profile: at initiation of heating, 26
25 deg.C; at 39 min., 90 deg.C; at 45 min., 100 deg.C; at 49 min., 104 deg.C; at 55 min., 106 deg.C; at 59 min, 105 deg.C, at which point the heat was removed and the reactor and contents cooled in an ambient temperature water bath having a temperature of between 20-30 deg.C.
30 When the reactor contents reached a temperature of 35 deg.C, the reactor was opened and the resulting coemulsion evaluated. The results are denoted on Table C1.

35

Example C20

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The coemulsification process of Example C19 was repeated with the same constituents used in forming Example C19, with the variation that additional ammonia was introduced to the initial constituent mixture in 5 order to evaluate the effects on the acidity of the polyethylene and its effects on the coemulsification process and the coemulsion. The resultant coemulsion was evaluated, and the results denoted on Table C1.

10

Example C21

Constituents comprising a low density polyethylene with an different acid number then previously utilized and listed on Table C1 were introduced into the reactor 15 which was sealed and the reactants stirred with the constituents remaining at 24-26 deg.C over a period of 15 minutes. The reactor was then heated in accordance with the following temperature profile: at initiation of heating, 26 deg.C; at 26 min., 101 deg.C; at 31 20 min., 104 deg.C; at 39 min., 104 deg.C; at which point the heat was removed and the reactor with contents cooled in an ambient temperature water bath having a temperature of between 20-30 deg.C. When the reactor contents reached a temperature of approximately 35 25 deg.C, the reactor was opened and the resulting coemulsion evaluated. The results are denoted on Table C1.

Example C22

30 A coemulsion was prepared in a two-step process where an intial "pre-coemulsion" material was produced, followed by subsequent formation of the coemulsion using the pre-coemulsion material.

35 Into the reactor was introduced a 128 grams of A- C 629 low molecular weight polyethylene having an acid

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value of 15.7, and 32 grams of Dow CSF amino siloxane. The reactor was sealed and the constituents stirred, during which the temperature of the constituents was raised to 112 deg.C, to assure the melting of the 5 polyethylene and thereafter maintained at that temperature for 5 minutes, subsequent to which the reactor and its contents were cooled. The reactor contents appeared as a hard waxy material of homogeneous nature, and was a beige color. This 10 material formed the pre-coemulsion material.

In the next step, to the pre-coemulsion material was added the remaining constituents as outlined on Table C1, and the reactor was then sealed, stirring was initiated, and the reactor was then heated in 15 accordance with the following temperature profile: at initiation of heating, 24-26 deg.C, after which the constituents were heated to 104 deg.C, at which point the constituents were retained in the temperature range of 104-115 deg.C for a time interval of 10 minutes. 20 Afterwards, heat was removed and the reactor was cooled in an ambient temperature water bath having a temperature of between 20-30 deg.C. When the reactor contents reached a temperature of approximately 35 deg.C, the reactor was opened and the resulting 25 coemulsion evaluated. The results are denoted on Table C1. The resultant coemulsion was somewhat cloudy in appearance, and was observed to contain particles.

Example C23

30 As in Example C22 a further coemulsion was prepared in a two-step process. The quantities of the materials used are denoted on Table C1

35 Into the reactor was introduced a quantity of A-C® 629 low molecular weight polyethylene having an acid value of 15.7, which was introduced onto the reactor

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which was then heated to 105 deg.C in order to melt the polyethylene, and once melted, the Dow CSF amino siloxane was poured onto the reactor. Immediate thickening of the mixture was noted, and subsequently

5 the mixture was poured from the reactor into an aluminum tray within which the mixture, which is the pre-coemulsion material, solidified.

In the next step, to the pre-coemulsion material was first broken down by crumbling into small pieces, 10 and introduced into the reactor, after which the remaining constituents as outlined on Table C1 were added. The reactor was then sealed, stirring was initiated, and the reactor was then heated in accordance with the following temperature profile: at

15 initiation of heating, 24-26 deg.C, after which the constituents were heated rapidly to 92 deg.C, then more slowly so that after 19 minutes the temperature rose to 104 deg.C, and after a further 10 minutes, the temperature rose only to 108 deg.C, thereby maintaining

20 the constituents for a time interval of 10 minutes in a temperature range between 104-108 deg.C. Afterwards, heat was removed and the reactor was cooled in an ambient temperature water bath having a temperature of between 20-30 deg.C. When the reactor contents were

25 sufficiently cooled, the reactor was opened and the resulting coemulsion evaluated. The results are denoted on Table C1. The resultant coemulsion was somewhat cloudy in appearance, and was observed to contain particles.

30

Example C24

To the reactor was added all of the individual constituents as outlined in Table C1, after which the reactor was sealed and the electric motor was energized 35 in order to drive the agitator. The contents of the

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reactor were then rapidly heated from a temperature of approximately 24 deg.C to 90 deg.C at a first rate of approximately 2 deg.C per minute, and then the heating rate was maintained at a reduced second rate of

5 approximately 1 deg.C per minute to raise the reactor contents from 90 deg.C to 104 deg.C Subsequently, the temperature was maintained in the range of 104 - 108 deg.C for a period of 10 minutes. Then, the heat was removed, and the contents cooled to a temperature of

10 approximately 30-35 deg.C. The agitator was then stopped, and the reactor was opened. The resultant emulsion was observed to produce a clear slide, and further tested with the results indicated on Table C1.

15

Example C25

As in Examples C22 and C23 a further coemulsion was prepared in a two-step process. The quantities of the materials used are denoted on Table C1

20 Into the reactor was introduced a quantity of A-C® 629 low molecular weight polyethylene having an acid value of 15.7, which was then heated to 125 deg.C in order to melt the polyethylene, and once melted, the potassium hydroxide (KOH) in the form of flakes of 87% concentration was commixed with water and added to the

25 reactor containing the polyethylene. After five minutes, to the reactor was further added the Dow CSF amino siloxane which was poured onto the reactor at which time it was observed that the constituents were thickening. The agitation was continued for a further

30 1-2 minutes, after which the agitator was halted and the constituents which form the pre-coemulsion material was poured into a try and allowed to solidify. The solidified material appeared to be waxy, and yellowish in color.

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In the next step, to the pre-coemulsion material was removed from the tray, and after being first broken down by crumbling into small pieces, reintroduced into the reactor, after which the remaining constituents as 5 outlined on Table C1 were added. The reactor was then sealed, stirring was initiated, and the reactor was then heated in accordance with the following 10 temperature profile: at initiation of heating, 24-26 deg.C, after which the constituents were heated rapidly to 76 deg.C, then more slowly at a rate of 1 deg.C/min. from 76 deg.C to 104 deg.C., at which time the heat was controlled so to maintain the temperature in the range 15 of 104-110 deg.C for 10 minutes. Afterwards, the heating means was removed and the reactor was cooled in an ambient temperature water bath having a temperature of between 20-30 deg.C. When the reactor contents were sufficiently cooled, the reactor was opened and the resulting coemulsion evaluated. The results are denoted on Table C1. The resultant coemulsion did 20 reveal a few large wax particles which were trapped on the surface and had not melted, however the balance of the coemulsion formed was observed to be only slightly cloudy in appearance, and otherwise no particles could be observed.

25

Examples K1-K13

Further samples having the overall compositions outlined in Table KS-1 were prepared in general accordance with the apparatus and the procedure outlined above for samples C10-C15.

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TABLE KS-1

Material:	K1	K2	K3	K4	K5	K6	K7	K8	K9	K10
A-C 629	32	32	32	32	32	32	32	32	34	34
A-C 329										
Dow CSF	8	8	8	8	8	8	8	8	6	6
Dow SSF										
Union Carbide Magnasoft	12.8	12.8	12.8	12.8	12.8	12.8	12	12	12.8	13
Ethomeen 18/12										
Igepal CO-430	3.2	3.2	3.2	3.2	3.2	3.2	4	4	3.2	3
Varonic T-202										
Tergitol 15-S-3	3.2	3.2	3.2	3.2	3.2	3.2	3.2	3.2	3.2	3
Ethal TDA-3	1.8	1.8	1.8	1.8	1.8	1.8	2.4	4.13	3.5	1.8
glacial acetic acid										2
hydrochloric acid, 10% sodium chloride										
sodium metabisulfite										
water	128.6	128.6	128.6	128.6	128.6	128.6	166	128.1	128.6	128.6
composition pH	4.87	5.03	4.9	4.8	4.97	4.77	4.51	4.77	5.1	5.05
Klett value:	185	290	310	330	345	365	465	460	295	250
optical density	0.37	0.58	0.62	0.66	0.69	0.73	0.93	0.92	0.59	0.5
light trans., %	42.7	26.31	24	21.9	20.43	18.64	11.76	12.05	25.74	31.64
slide appearance	clear	clear	clear	cloudy	cloudy	cloudy	clear	clear	clear	clear

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K11	K12	K13
====	====	====

34	34	34
----	----	----

6	6	6
---	---	---

12	12	12
4		4

3.5	3.5	3.5
-----	-----	-----

0.4	0.4	0.4
-----	-----	-----

128.1	128.1	128.1
-------	-------	-------

5.15	5.1	5.12
------	-----	------

350	370	395
0.7	0.74	0.79
20.93	18.2	16.23
clear	clear	clear

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The concentration of the materials were varied in the percentages outlined. Results listing the acidity and the optical qualities of the sample are listed.

5

Examples P1-P8

In order to determine the dynamic coefficient of friction, or "COF" of the inventive compositions as used with a paper substrate, compositions designated P1-P8 as outlined in Table PS-1 were produced in 10 accordance with the general methods used for the production of examples C1-C15.

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Material:	P1	P2	P3	P4	P5	P6	P7	P8
A-C 629					34	32		176
A-C 392	34	32						
Dow CSF	6	8	6	8				
Dow SSF								
Union Carbide Magnasoft					12	12.8		
Ethomeen 18/12	12	12					44	44
Igepal CO-710								
Igepal CO-430	4	4						
Varonic T-202								
Tergitol 15-S-3					3	3.2		
Ethal TDA-3	3.5	3.5	2	1.8				
glacial acetic acid								
hydrochloric acid, 10%	0.4	0.4				1.76		
sodium chloride							3.78	3.78
sodium metabisulfite							224.4	528
potassium hydroxide								533
water	128.1	128.1	128.6	128.6				
composition pH	5.05	5.1	5.1	5.2	5.2	9.2	9.5	
Klett value:	295	290	240	224	55		210	
optical density	0.59	0.6	0.48	0.44	0.11	0.42		
light trans., %	25.7	25.1	33.15	35.6	77.6	38		
slide appearance	clear	clear	clear	clear	clear	clear		

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For comparative purposes, four compositions were formulated as control samples for the purpose of comparison. These included: sample P5 which was a amino silicon control; sample P6 which comprised low density polyethylene; sample P7 which comprised high density polyethylene; and sample P8 which was a paper control blank having no lubricant coating. The paper substrate was Westvaco Sterling II Litho Gloss, a web offset paper having an 80# weight rating. Compositions P1 - P8 were all coated onto the paper using a wire wound #6 draw down rod, which gave a 0.54 mil wet film thickness. All of the compositions were applied at a 30 % non-volatile level to ensure consistent and equal film deposition.

The evaluation of each of the compositions was performed under two different contact situations: "coated/coated" and "coated/uncoated". What is meant by "coated/coated" is two contacting faces of the paper substrate used in any particular evaluation were coated with a particular composition so that a measure of the dynamic coefficient of friction of the two coated faces could be evaluated. In a similar manner, what is meant by "coated/uncoated" is that one of two contacting faces of the paper substrate used in an evaluation was coated with a composition so to provide a measure of the dynamic coefficient of friction of one coated face against a non-coated or "neat" face of paper could be determined. The coefficient of friction was determined by the use of a "Coefficient of Friction Tester" Model No. 32-25 produced by Testing Machines, Inc. This device includes an inclining board with a weighted sled and measures the angle, at which the individual coated substrates allowed the weighted sled to move. The tangent value of this angle is used as the COF for any test.

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Test results for each of the compositions P1 - P8 were determined for both "coated/coated" and "coated/uncoated" evaluations from four individual testing trials of each. The results from each of the 5 four trials for a particular composition were averaged to determine the dynamic COF values the results of which are reported in Table PS-2 below.

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TABLE PS-2
Example Composition:

	Coefficients of Friction (Dynamic)	
	Coated/Coated	Coated/Uncoated
P2	0.2378	0.1899
P3	0.2171	0.1944
P4	0.275	0.2633
P5	0.2773	0.2633
P6	0.2726	0.2424
P7	0.4296	0.4117
P8	0.2656	0.2679
PC1	(Uncoated/Uncoated) = 0.337	

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As may be seen from these test results, the resulting COF values as determined for the coemulsion compositions, whether such coemulsions utilized low density or high density polyethylenes in their 5 formulations provided generally improved values in the reported performance characteristics as compared to compositions of emulsions having only a single constituent. This may be particularly seen in conjunction with the results of associated with 10 compositions P2 and P3.

The compositions were tested in conjunction with fiber and fabric (textile) samples in order to evaluate the effects of the use of the compositions. Fabric samples were evaluated by treating samples of a 65/35 15 polyester/cotton filter cloth where each of the fabric samples was a panel of dimensions 14.5 inches by 18 inches. The fabric samples were prepared by separately treating each within a fluid bath which contained one of the compositions labeled as T1 - T16 whose 20 compositions are described on Table TS-1 below.

-51-

Material:	T1	T2	T3	T4	T5	T6	T7	T8
A-C 629				34	32	34	32	176
A-C 392				34	32		176	
Dow CSF				6	8	6	8	
Dow SSF								
Union Carbide Magnasoft								
Ethomeen T-12				12	12	12	12.8	
Ethomeen 18/12								
Igepal CO-710				4	4			
Igepal CO-430								
Varonic T-202								
Tergitol 15-S-3								
Ethal TDA-3						3	3.2	
glacial acetic acid				3.5	3.5	2	1.8	
hydrochloric acid, 10%								
sodium chloride				0.4	0.4			
sodium metabisulfite								
potassium hydroxide								
water				128.1	128.1	128.6	128.6	124.4
composition pH					5.05	5.1	5.2	9.2
Klett value:				295	290	240	224	55
optical density				0.59	0.6	0.48	0.44	0.11
light trans., %				25.7	25.1	33.15	35.6	77.6
slide appearance				clear	clear	clear	clear	clear

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	T9	T10	T11	T12	T13	T14	T15	T16
	34	32		34	32			176
	6	8	6	6	8			
159							160	
	12	12		12	12.8			
	4	4				44	44	
	3.5	3.5		3	3.2			
	0.4	0.4		2	1.8			
						1.76		
530	128.1	128.1		128.6	128.6		3.78	
	5.05	5.1	5.1		5.2		528	
	295	290	240		224	55	210	
	0.59	0.6	0.48		0.44	0.11	0.42	
	25.7	25.1	33.15		35.6	77.6	38	
	clear	clear	clear		clear	clear	clear	

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These fluid baths are well known to the textile processing art, and typically comprise a major proportion of water, and minor proportions of other additives such as softening agents, various resins, 5 including permanent press resins, as well as other desired constituents which are useful in the treatment of textiles.

With regard to these samples, sample T1 was a control sample consisting of water with no further 10 additives, samples T2-T5 were samples comprising coemulsions according to the invention and amino silicone, samples T6-T8 were additional comparative samples of various emulsions, sample T9 was a comparative example which included a blend of water and 15 a durable press resin, "Reactex LFF" which is described as a self-catalyzed glyoxal based material which is used as a permanent press finishing agent and features low formaldehyde shrinkage control, which is commercially available from Ivax Industries, samples 20 T10-T13 were samples comprising coemulsions according to the invention and amino silicone, and further included the durable press resin, samples T14-T16 were further examples with various emulsions. The emulsions were prepared in a manner similar to the methods used 25 for the production of examples which are outlined on Table C1 noted above. The individual samples were contacted to the fabric by methods known to the art, so that the fabric had a 56% wet pick-up of the fluid and each of the individual samples were evaluated for the 30 following physical characteristics: tensile strength according to the AATCC test protocol, "hand" using a four point scale, wherein a value of 1 is indicative of superior "hand" characteristics, and a value of 4 is indicative of poor "hand" characteristics, "Hunter 35 whiteness", wrinkle recovery according to the AATCC

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test protocol, the Stoll flex in both the warp and fill directions of the fabric samples, and the tear strength in both the warp and fill directions of the fabric samples. The results of the testing of the fabric samples utilizing the compositions described on Table TS-1 is outlined on Table TS-2 below.

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TABLE TS-2

Sample:	Tensile Str. in lbs.	Hand Whiteness	Hunter Whiteness	Wrinkle recovery	Stoll Flex (warp/fill)	Tear Strength (warp/fill)	Needleburn
T1	44	2	126	264	625/375	3.4/2.6	clogged
T2	41	1	123	274	3750/2450	5.8/4.4	clean
T3	40	1	124	272	6350/3350	5.7/4.4	clean
T4	42	1	123	270	---	5.4/4.3	clean
T5	42	1	123	272	5700/3300	5.3/4.3	clean
T6	42	1	124	278	5000/4300	5.6/4.1	clean
T7	42	2	122	265	2350/1150	4.9/4.2	clean
T8	38	2	123	286	1250/1150	6.3/5.1	clean
T9	47	—	128	294	1250/325	3.4/2.1	partial clog
T10	44	—	123	293	5600/3100	5.1/3.6	clean
T11	42	—	125	302	6500/3100	5.1/3.7	clean
T12	43	—	124	299	4850/2650	4.6/3.3	clean
T13	42	—	124	—	3500/1650	4.6/3.1	clean
T14	40	—	128	304	6500/2850	5.0/3.9	clean
T15	44	—	128	289	2700/2050	4.7/3.2	clean
T16	40	—	124	307	2050/1650	5.5/4.1	clean

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As may be seen from the results on Table TS-2 the compositions utilizing the coemulsions all show tensile strength properties, Hunter Whiteness values, and wrinkle recovery values which are comparable to the 5 samples of a single constituent emulsion, namely T6-T8, T14-T16. A striking improvement in the hand is to be noted for compositions T2-T6 which comprise the coemulsions, as well as modest to marked improvements 10 in the Stoll Flex characteristics of the fabric samples treated with the coemulsions as compared to the results observed for emulsions containing a single constituent, namely T6-T8, and T14-T16.

The compositions denoted as T1-T16 were also evaluated in conjunction with a fabric panels of 15 65%/35% polyester/cotton fiber cloth in order to evaluate the needleburn characteristics of the treated fabric panels. The fabric panels were prepared in accordance with the fabric panels listed on Table TS-1. The lubricated fabric panels were tested using 16/1 20 needles on a Singer model 282 sewing machine operating at a rate of 5000 stitches per minute running through 4 folds of fabric through 4 passes. The observed characteristics from this testing procedure are noted on Table TS-2. As may be seen from these results, the 25 needle remained clean through all of the samples except for the two control compositions, T1 and T9 which contained no emulsion.

It can be seen by the foregoing that the coemulsions according to the instant invention provide 30 compositions useful as processing aid for improving fabric, thread and paper processing characteristics, particularly in improving the "hand", of textiles treated with the coemulsions taught herein, which provide such improved characteristics with relatively 35 small amounts of amino functional silicone softeners in

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conjunction with other constituents. Such an advance allows for a more cost efficient use of such silicone containing constituents with no compromise in performance characteristics, but rather a surprising

5 improvement in key properties of the materials with which the coemulsion is used.

It will be appreciated that the instant specification and the examples set forth are by way of illustration and not by limitations, and that various 10 modifications and changes may be made to the teachings presented herein, and that various modifications and changes may be made without departing from the spirit and scope of the present invention and are to be considered a part thereof, said invention being limited 15 only by the following claims.

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CLAIMS

1. An aqueous composition comprising:
a coemulsion which includes;
at least one oxidized polyethylene selected from the group
consisting of: polyethylene homopolymers including high
05 density polyethylene homopolymers, low density polyethylene
homopolymers, polyethylene/acrylate copolymers, polyethylene
terpolymers containing acrylates, and polyethylene terpolymers
containing esters, and
10 at least one amino functional silicone.
2. The article of claim 1 wherein the amino functional
silicone is a polydiorganosiloxane.
15
3. The article of claim 2 wherein the polyorganosiloxane
has an amine neutral equivalent in the range of between about
1000 and 3000.
20
4. The aqueous composition of claim 1 wherein the
composition further comprises:
at least one constituent selected from the group consisting
of: ethoxylated aliphatic amines,
25 ethoxylated octylphenols,
ethoxylated nonylphenols,
ethoxylated primary alcohols,
ethoxylated secondary alcohols,
nonionic emulsifiers,
30 fatty acids,
ammonium hydroxide, and
sodium salts.
5. The composition of claim 4 wherein the ethoxylated
35 aliphatic amine are the reaction products of a saturated or
unsaturated fatty amines reacted with ethylene oxide.

6. A process for producing an aqueous composition comprising the coemulsion according to any preceding claim which process includes the steps of:

providing the oxidized polyethylene and the amino functional silicone to a reaction vessel,
raising the temperature of the oxidized polyethylene and amino functional silicone to at least the melt point of the oxidized polyethylene constituent.

10 7. The process according to claim 6 which further includes the process steps of:

raising the temperature of the constituents beyond the melt point of the oxidized polyethylene constituent to a maximum temperature,

15 followed by maintaining the constituents at this maximum temperature for a first residence time interval.

8. The process according to claim 6 which includes the process steps of:

20 raising the temperature of the oxidized polyethylene and amino functional silicone, with agitation, at a first heating rate through the melting point of the oxidized polyethylene component for a first time interval,

25 then during a second time interval, further heating at a second heating rate wherein the first heating rate is greater than the second heating rate to raise the temperature to a maximum temperature or maximum temperature range approximately 7-10 deg.C beyond the melt point of the oxidized polyethylene component, and,

30 maintaining the reactants at this maximum elevated temperature or maximum temperature range for a period of between about 10-15 minutes.

9. The process according to claim 6 which includes a further process step of adjusting the pH of the coemulsion to maintain the pH in the range of 4-6.

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10. An article comprising at least a partial coating of the composition according to any of claims 1 - 5 which article is selected from the group which includes: textiles, fibers, ropes, tows, webs, threads, fiberfill material, paper, release sheets, release strips, asphalt comprising articles such as roofing paper and roofing shingles.

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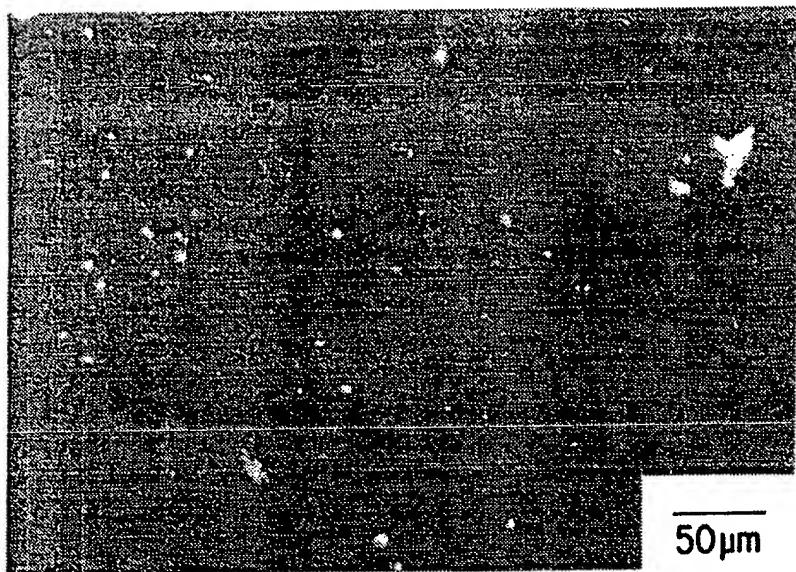


FIG. 1

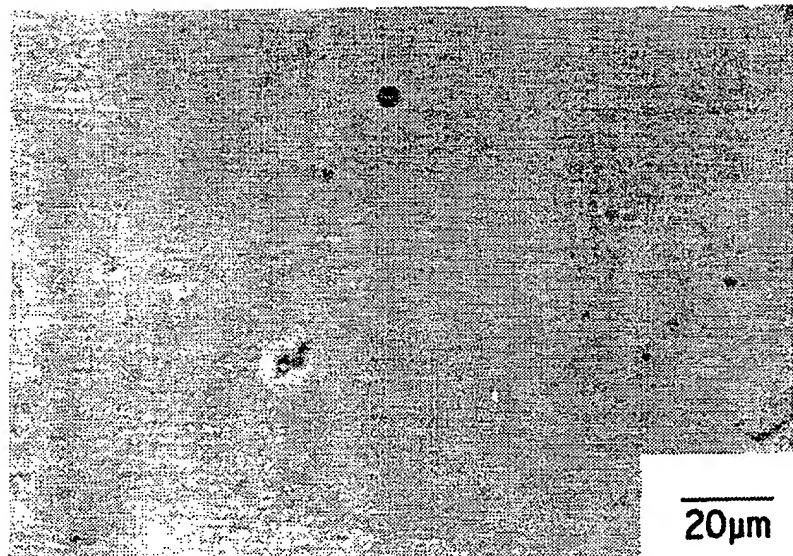


FIG. 2

2/4

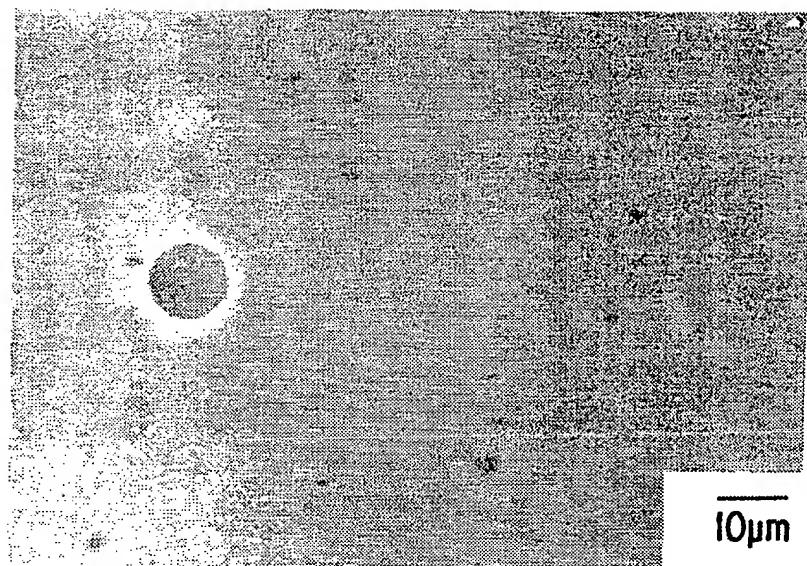


FIG. 3



FIG. 4

3/4

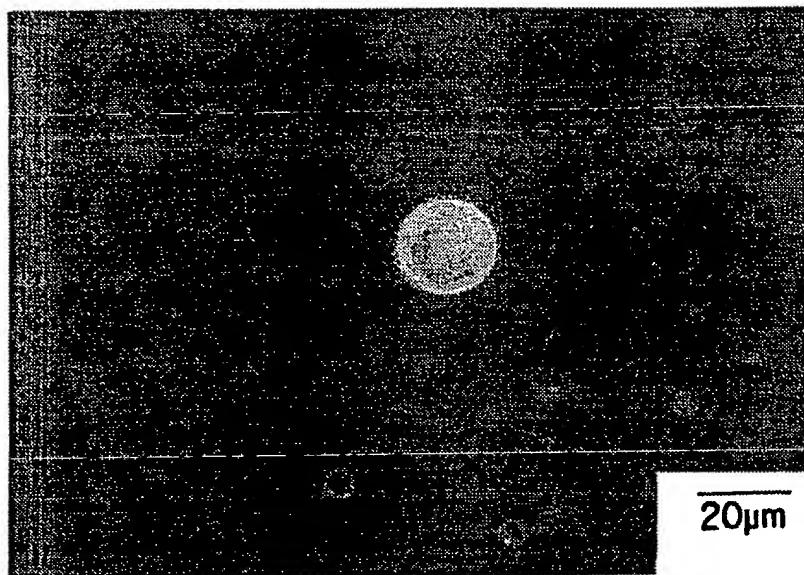


FIG. 5

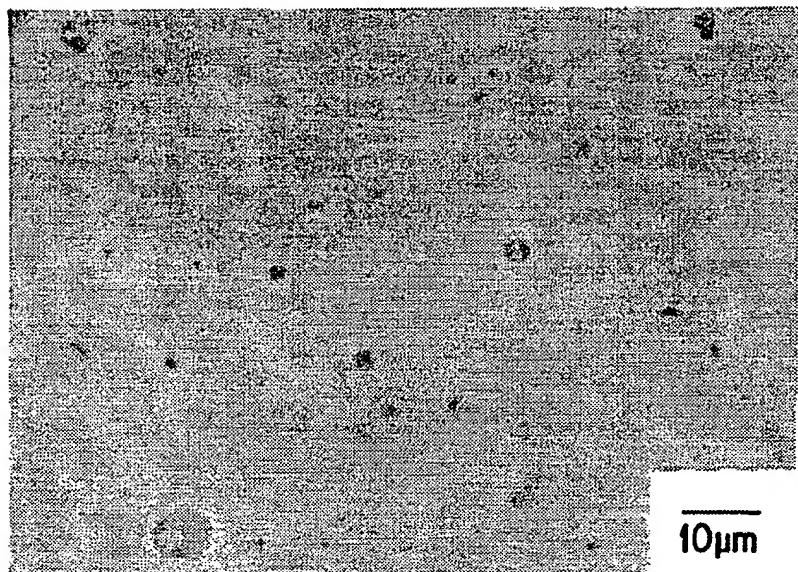


FIG 6

4/4

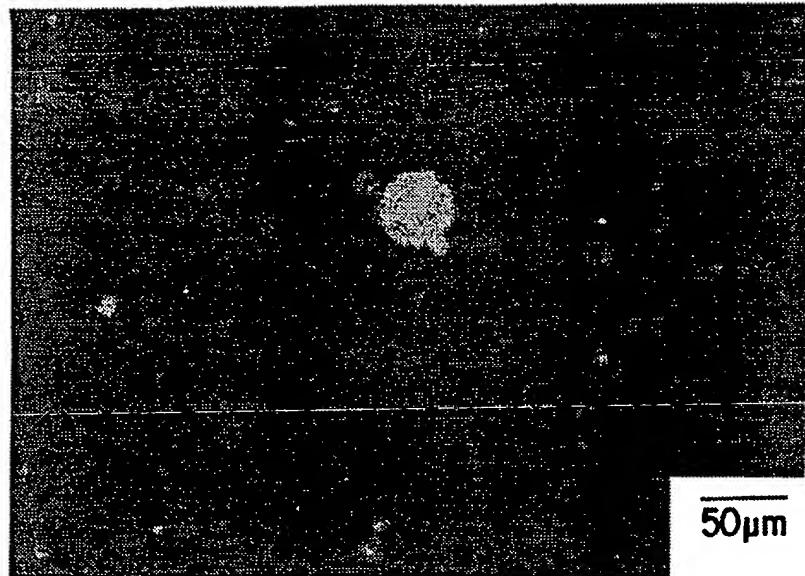


FIG. 7

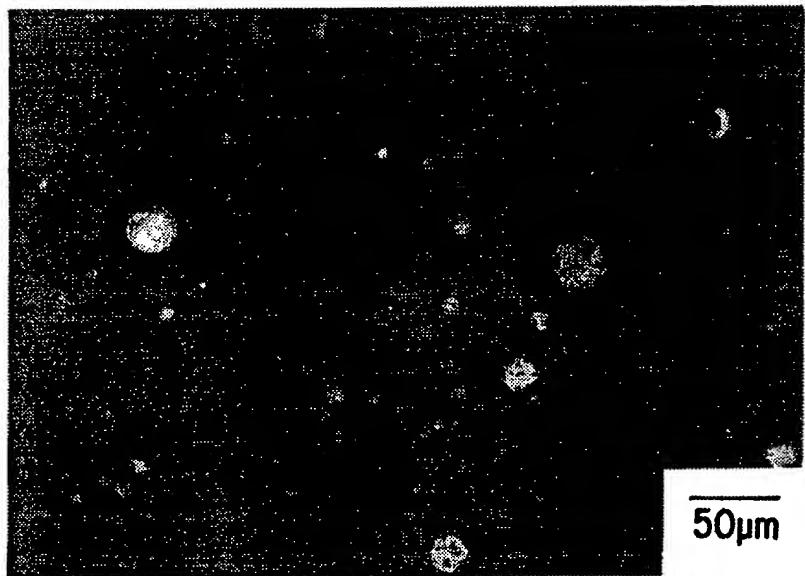


FIG 8

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 91/06342

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all)⁹

According to International Patent Classification (IPC) or to both National Classification and IPC

Int.C1.5 C 08 L 23/06 D 06 M 15/21 C 09 J 7/02

II. FIELDS SEARCHED

Minimum Documentation Searched⁷

Classification System	Classification Symbols		
Int.C1.5	C 08 L	C 08 J	C 09 D
	C 09 J	D 06 M	

Documentation Searched other than Minimum Documentation
to the Extent that such Documents are Included in the Fields Searched⁸

III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹

Category ⁹	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
P, X	EP, A, 0412324 (CHEM. FABR. PFERSEE) 13 February 1991, see claim 1 ---	1-10
X	EP, A, 0220400 (ALLIED CORP.) 6 May 1987, see claim 1, & US, A, 4767646 (cited in the application) ---	1-10
Y	EP, A, 0189978 (UNITED MERCHANTS & MANUFACTURERS INC.) 6 August 1986, see claims 1, 3, 7 ---	1-10
Y	EP, A, 0244500 (MITSUBISHI CHEM.) 11 November 1987, see page 1, lines 32, 36-37; claim 3 ---	1-10
Y	US, A, 3904805 (JOHNSON, BERUBE) 9 September 1975, see claims 1, 6 ---	1-10

⁹ Special categories of cited documents : ¹⁰

- ^{"A"} document defining the general state of the art which is not considered to be of particular relevance
- ^{"E"} earlier document but published on or after the international filing date
- ^{"L"} document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- ^{"O"} document referring to an oral disclosure, use, exhibition or other means
- ^{"P"} document published prior to the international filing date but later than the priority date claimed

- ^{"T"} later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- ^{"X"} document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step
- ^{"Y"} document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- ^{"&"} document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

14-01-1992

Date of Mailing of this International Search Report

05.02.92

International Searching Authority

EUROPEAN PATENT OFFICE

Signature of Authorized Officer

MORTENSEN

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No.
A	EP,A,0291213 (TORAY) 17 November 1988, see page 3, lines 12-15; claim 1 ----	1
A	EP,A,0230565 (HOFFMANN'S STÄRKEFABRIKEN AG) 5 August 1987, see claim 1 ----	1
A	WO,A,8912549 (AVERY INT.) 28 December 1989, see page 11, lines 31-37; claim 1 -----	1

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

US 9106342
SA 52221

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 30/01/92. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
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		JP-A-	3076735	02-04-91
EP-A- 0220400	06-05-87	US-A-	4767646	30-08-88
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EP-A- 0189978	06-08-86	AU-A-	5217586	17-07-86
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		AU-B-	598492	28-06-90
		AU-A-	6584886	04-06-87
		WO-A-	8703682	18-06-87
		JP-T-	63502362	08-09-88
		US-A-	4818242	04-04-89
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		EP-A-	0379561	01-08-90
		JP-T-	3500060	10-01-91